

WO9957201

Publication Title:

Compositions comprising a thermoplastic component and superabsorbent polymer

Abstract:

The present invention relates to a composition comprising a thermoplastic component and at least one superabsorbent polymer, methods of use for such composition, and articles comprising such composition. The composition may be formed into a film layer or applied to an article with various hot melt adhesive application techniques. The composition is useful for a variety of end-uses, particularly for use in disposable absorbent articles such as disposable diapers, feminine napkins and medical dressings.

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C08L 101/00, A61L 15/00, H01B 7/28	A1	(11) International Publication Number: WO 99/57201 (43) International Publication Date: 11 November 1999 (11.11.99)
(21) International Application Number: PCT/US99/10135 (22) International Filing Date: 7 May 1999 (07.05.99) (30) Priority Data: 60/084,582 7 May 1998 (07.05.98) US 09/306,265 6 May 1999 (06.05.99) US (71) Applicant (for all designated States except US): H.B. FULLER LICENSING & FINANCING, INC. [US/US]; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): AHMED, Sharf, U. [-/US]; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US). EMIRU, Andualem, W. [-/US]; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US). CLAPP, Leslie, J. [-/US]; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US). KROLL, Mark, S. [-/US]; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US). (74) Agents: QUAN, Nancy, N. et al.; 1200 Willow Lake Boulevard, Box 64683, St. Paul, MN 55164-0683 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: COMPOSITIONS COMPRISING A THERMOPLASTIC COMPONENT AND SUPERABSORBENT POLYMER		
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COMPOSITIONS COMPRISING A THERMOPLASTIC COMPONENT AND SUPERABSORBENT POLYMER

5 **Related Applications**

This application is a Continuation-In-Part of U.S. Patent Application Serial No. 60/084,582 filed May 7, 1998.

Field of the Invention

10 The present invention relates to a composition comprising a thermoplastic component and at least one superabsorbent polymer, methods of use for such composition, and articles comprising such composition. The composition may be formed into a film layer or applied to an article with various hot melt adhesive application techniques. The composition is useful for a variety of end-uses, particularly
15 for use in disposable absorbent articles such as disposable diapers, feminine napkins and medical dressings.

Background of the Invention

Water-insoluble, water swellable hydrogel-forming absorbent polymers, also
20 referred to as superabsorbent polymers (SAP), are capable of absorbing large quantities of liquids such as water, body fluids (e.g., urine, blood), industrial fluids, household fluids and are further capable of retaining such absorbed liquids under moderate pressures.

For example, SAP is employed in various cable applications to shield the
25 penetration of moisture. When the outer sheath of the cable is damaged allowing penetration of fluids, the SAP absorbs the incoming liquid and swells, forming a physical barrier to the entry of any further fluid. Particulate SAP is typically held in place by means of a hydrosoluble binder. For example, U.S. Patent No. 4,837,077 issued to Anton et al., June 6, 1989, relates to a hydroexpansible composite material including a
30 solid flat support having on at least one of its faces a coating with a high proportion of a hydroexpansible polymer powder fixed to the support by means of a hydrosoluble binder, the coating being essentially formed of 95% to 55% by weight of a hydroexpansible

polymer powder, 5% to 23% by weight of a hydrosoluble binder, and surfactant in a positive amount of up to 22% by weight. The ingredients are homogenized by means of a liquid. This reference expressly teaches that thermoplastic substances cause an unfavorable effect, the blockage of a part of the hydroexpansible polymer powder, thus
5 limiting its ability to swell in the presence of water.

The absorption characteristics of SAP materials make them especially useful for incorporation into absorbent articles such as disposable diapers, adult incontinent pads and briefs, and catamenial products such as sanitary napkins and the like. A desired characteristic of such disposable absorbent articles is thinness. Thinner disposable
10 articles are less bulky to wear, better fitting under clothing and are less noticeable. Further, thin disposable absorbent articles are more compact, making the product easier for the consumer to carry and store. Compactness in packaging also results in reduced distribution costs for the manufacturer and the distributor, including less shelf space. The ability to provide thinner absorbent articles such as a diaper has been contingent on
15 the ability to develop relatively thin absorbent cores or structures that can acquire and store large quantities of fluid, particularly urine. Therefore, there is a trend towards employing higher concentrations of SAP polymers to achieve this purpose.

SAP is typically available in a particulate or powder form that is sprinkled or sifted into the absorbent core portion, which is sandwiched between a fluid pervious
20 topsheet and a fluid impervious backsheet. The incorporation of particulate SAP into an article tends to generate dust from the SAP fines. Further, conventional absorbent articles have the limitation that the absorbent gelling particles are not immobilized and are free to migrate and shift during the manufacturing process and/or use. Movement of the SAP particles during manufacture can lead to absorbent material handling losses
25 during manufacturing operations as well as improper distribution of the particles.

Other problems occur when these absorbent gelling particles migrate during and after swelling. The inability to fix the particles at optimum locations leads to insufficient fluid storage in one area and over-capacity in other areas due to the movement of the SAP particles to locations other than where originally positioned. Another important
30 factor that has to be considered is the liquid permeability of the SAP. It has been discovered that the fluid transport properties of the gel layer formed as a result of the

swelling SAP particles in the presence of fluids is extremely important. Although the formation of a SAP gel layer fluid barrier, known as "gel blocking" is desirable for some applications, such as for use in cables, the formation of gel layers in disposable absorbent products is undesirable since it greatly reduces the efficiency of the SAP. Thus, the advantages of being able to fixate SAP in place are apparent and several ways of accomplishing such have been suggested.

For example, U.S. Patent No. 4,392,908 issued to Dehnel July 12, 1983, concerns a process for manufacturing a water-absorbent article in which particles of a water swellable polymer are fixed to a water-absorbent substrate. The process includes the steps of forming on the surface of water-swallowable particles a coating of a thermoplastic adhesive resin; locating the coated particles in their unswollen state and dry state on or within the water-absorbent substrate also in the dry state; and applying heat to soften the thermoplastic coating of the particles and pressing the particles and substrate to cause the particles to be bound to the substrate. The particles of absorbent material can be coated by mixing the absorbent particle with an aqueous latex of the resin and then drying the mixture, spraying a resin solution or latex onto the particles, adding the resin to the particles when the particles are in a slurry, or incorporating the thermoplastic as a solution or latex during the manufacture of the absorbent material.

Further, SAP particles may be combined with certain thermoplastic compositions. U.S. Patent No. 4,977,211 issued to Doi, Dec. 11, 1990, teaches a moldable water-absorptive resin composition which comprises (A) 5% to 95% by weight of a water-absorptive resin and 95% to 5% by weight of (B) a polyolefin resin and (C) an ethylene/alpha-olefin copolymer rubber based on the total content of said components (A), (B) and (C), wherein the contents of (C) amounts to 20% to 80% by weight based on the total contents of said component (B) and (C). An ethylene/vinyl acetate copolymer is exemplified having a vinyl acetate content of 33% by weight and a melt index of 30 g/10 min. Subsequently, the water-absorptive resin composition of the present invention can be molded into a form suitable for its final use. The molded products may be subjected to a secondary molding treatment such as foaming or orientation. The compositions can be molded alone into a film, a sheet, fibers or other products.

Japanese Patent Application 9-176618 published July 8, 1997 relates to a water-swelling hot melt pre sealant for lap joint construction external wall material. The composition includes 20 wt-% to 50 wt-% thermoplastic elastomer, 5 wt-% to 40 wt-% tackifier, 10 wt-% to 50 wt-% plasticizer and 5 wt-% to 35 wt-% water swellable resin.

5 The plasticizer is described as paraffin type oil, naphthene type oil, aromatic type oil, polybutene liquid rubber, polyisobutylene liquid rubber, polyisoprene liquid rubber and its hydrogenated product.

Japanese Patent Application 60-195299, published Oct. 3, 1985 pertains to a method for preventing water leaks with good operability, especially in engineering work

10 and construction employing a water-swellaable composition styrene elastomers, highly water absorbing materials and tackifiers. The proportions of the ingredients are usually 5-100 parts of the highly water-absorbing resin and 100 to 200 parts of tackifier based on 100 parts of the styrene elastomer.

More recently, WO 98/27559A1, published June 25, 1998 concerns a water

15 swellaable thermoplastic compound for gluing and coating made from a water insoluble component, a water soluble or water dispersible component, and a water-swellaable component. Examples of suitable water-soluble or water dispersible oligomers, homopolymer or copolymers include polyethylene glycol in the molecular weight range between 2400 and 20,000, polyvinyl methyl ether, polyvinylpyrrolidone, copolymers of

20 vinyl methyl ether or vinyl pyrrolidone, polyvinyl alcohols, water soluble or water dispersible polyesters or copolyesters, and water soluble or water dispersible acrylate polymers.

Summary of the Invention

25 The present inventors have discovered that the inclusion of water soluble or dispersible polymers disadvantageously reduces the rate of fluid acquisition. It is surmised that water soluble/dispersible polymers compete with the SAP in absorption of the fluid. In such instances, the water soluble or dispersible ingredient forms a solution or dispersion that thickens the fluid intended to be absorbed by the SAP. The thickened

30 fluid tends to be absorbed by the SAP at a much slower rate than that of the fluid prior to thickening, thus hindering the rate of absorption.

Surprisingly, the present applicants have found that the fluid acquisition rate and the total absorption capacity for small particle size SAP is not impaired by being admixed with certain thermoplastic compositions. Further, in many instances the presence of the thermoplastic component actually enhances the performance of the SAP, particularly for reducing unintended gel blocking.

The present invention is a composition comprising a thermoplastic composition, also referred to as the "thermoplastic component" and at least one superabsorbent polymer. The thermoplastic component is sufficiently polar and simultaneously of relatively low cohesive strength. The cohesive strength corresponds to the melt index of a polymer or molten viscosity of a mixture of a thermoplastic polymer and other ingredients.

In one embodiment, the present invention is a thermoplastic composition comprising a thermoplastic component comprising at least one thermoplastic polymer and at least one water insoluble diluent having polar functionality and at least one superabsorbent polymer. The thermoplastic component is preferably water insoluble. The diluent is preferably a plasticizer employed at a concentration ranging from about 10 wt-% to about 30 wt-% of the total weight of the composition.

In another embodiment, the present invention is a thermoplastic composition comprising a thermoplastic component comprising at least one interpolymer of ethylene. The interpolymer has a melt index greater than about 500 g/10 min., preferably about 800 g/10 min. and greater, and more preferably about 1000 g/10 min. and greater. Further, the interpolymer of ethylene preferably has a comonomer content of greater than 25 wt-% and preferably greater than about 28 wt-% or higher. The comonomer is preferably vinyl acetate, methacrylate, n-butyl acrylate and mixtures thereof. Further, the composition is preferably free of water soluble/dispersible polymers having a molecular weight (Mw) of greater than 2000.

In another embodiment, the present invention is a thermoplastic composition comprising about 25 wt-% to about 50 wt-% of a thermoplastic component having a viscosity of less than about 30,000 cPs at 177°C and 50 wt-% to about 75 wt-% of a superabsorbent polymer.

The thermoplastic component may be a single thermoplastic polymer or a blend of thermoplastic polymers in its entirety. Alternatively, the thermoplastic component may comprise at least one thermoplastic polymer in combination with at least one hot melt adhesive ingredient such as plasticizers, tackifiers and waxes. The thermoplastic component is preferably present in an amount ranging from about 40 wt-% to about 70 wt-%, whereas the superabsorbent polymer is preferably present in an amount ranging from about 30 wt-% to about 60 wt-%. Further, the thermoplastic component preferably comprises at least one thermoplastic polymer including block copolymers, amorphous polyolefins, crystalline polyolefins, interpolymers of ethylene and mixtures thereof.

The composition preferably gels at a rate equal to or faster than that of the SAP polymer alone, particularly when employing SAP having a particle size of less than about 200 microns. The rate of gelation is preferably less than 3 hours, more preferably less than 1 hour and most preferably less than about 30 minutes.

The resulting composition can be used in place of particulate or powdered SAP for any application employing such, avoiding the aforementioned problems. The composition can advantageously be applied to an article by any known hot melt application technique, extruded as a rod, strand or fiber, as well as be formed into a film layer on a substrate, thus providing SAP in a roll good form.

In another embodiment, the invention is film layer disposed on a substrate comprising the inventive thermoplastic mixture. The film layer may be formed by providing a molten mixture of the inventive composition comprising at least one thermoplastic polymer and at least one superabsorbent polymer and forming a substantially continuous film from said molten mixture onto a substrate.

In another embodiment, the invention is a method of applying superabsorbent polymer to a substrate comprising the steps of:

- a) providing a molten mixture comprising a thermoplastic composition and at least one superabsorbent polymer; and
- b) applying said mixture to a substrate.

Detailed Description of the Invention

The term "thermoplastic composition" and "thermoplastic component" refers to a composition that is reversibly capable of softening or fusing when heated and hardening again when cooled.

5 The term "molten" as used herein refers to a flowable or liquified state of matter attained with heat.

The terms "water responsive" and "water sensitive" as used herein refers to ingredients that are soluble or dispersible in an aqueous environment. Aqueous environment includes neutral, basic, acidic and ionic aqueous solutions.

10 The term "water insensitive" as used herein refers to an ingredient that is soluble in water at a concentration less than 1.0 wt-%, and preferably less than 0.5 wt-%.

The present invention is a composition comprising a thermoplastic composition and at least one super absorbent polymer (SAP). The thermoplastic component provides the thermoplasticity of the SAP containing composition, whereas the SAP polymer
15 provides the fluid absorption capability of the composition. The thermoplasticity allows for a variety of application techniques, such as those employed for hot melt adhesives. The thermoplastic component also solves the gel blocking problem. Gel blocking is the phenomena wherein SAP particles absorb fluid and swell into a gel layer. The gel layer creates a barrier and prevents the migration of fluid to the ungelled SAP particles. Since
20 the thermoplastic component provides a means of separating the individual small particle size SAP particles from each other, each SAP particle can absorb fluid and swell independently, essentially unaffected by neighboring SAP particles. Hence gel blocking is reduced, increasing the rate of absorption of the SAP.

The thermoplastic component is sufficiently polar and simultaneously of
25 relatively low cohesive strength. Both these properties are contributed to by the selection of ingredients employed. The thermoplastic component comprises at least one thermoplastic polymer and at least one water insoluble diluent having polar functionality or at least one interpolymers of ethylene having a melt index greater than about 500 g/10 min and a comonomer content greater than about 25 wt-%, or a thermoplastic
30 composition having a viscosity of less than about 30,000 cPs at 350°F (177°C) in combination with at least 50 wt-% superabsorbent polymer. The thermoplastic

component is present at a concentration sufficient to form a continuous phase within the mixture. Hence, the concentration of thermoplastic component is dependent on the concentration and particle size of the superabsorbent polymer employed. In general, the thermoplastic component comprises from about 20 wt-% to about 80 wt-%, preferably
5 from about 30 wt-% to about 70 wt-%, and most preferably from about 40 wt-% to about 60 wt-%.

The thermoplastic composition may comprise, in its entirety, a single thermoplastic polymer, a blend of thermoplastic polymers, or alternatively the thermoplastic composition may be a hot melt adhesive comprising at least one
10 thermoplastic polymer in combination with other thermoplastic diluents such as tackifying resins and plasticizers. The invention also contemplates the thermoplastic composition being comprised entirely of at least one thermoplastic tackifier, plasticizer, wax and mixtures thereof. However, in most instances, a thermoplastic polymer is necessary to provide the desired properties, particularly strength. The thermoplastic
15 polymer alone may exhibit sufficient adhesive characteristics for some applications. However, preferably the thermoplastic polymer is combined with tackifiers and plasticizers to modify the adhesive properties for use in the intended application. This embodiment is particularly useful wherein the composition serves a dual purpose of providing the SAP in a non-particulate form as well as fixating the SAP in place. SAP
20 containing hot melt adhesive may also serve the function of bonding at least one substrate layer to at least one other substrate layer.

The thermoplastic component of the present invention typically comprises at least one thermoplastic polymer present in an amount ranging from about 10 wt-% to about 60 wt-%, preferably in an amount ranging from about 15 wt-% to about 50 wt-%, and more
25 preferably in an amount ranging from about 20 wt-% to about 50 wt-%, with respect to the total weight of the composition.

For embodiments that employ only a single thermoplastic polymer as the entire thermoplastic component, the thermoplastic polymer will typically either be sufficiently polar and/or combined with a polar diluent. In the case of employing blends of
30 thermoplastic polymers as the entire thermoplastic component, the blend will exhibit the same desired properties. For some embodiments, a relatively high viscosity

thermoplastic polymer is combined with relatively low viscosity thermoplastic components to enhance the cohesive strength of the mixture while maintaining good processability. For this embodiment, the relatively high viscosity polymer generally has a melt index (MI) of about 400 g/10 min. or less preferably about 200 g/10 min. or less, more preferably about 100 g/10min. or less and most preferably less than about 50 g/10 min. In order to achieve the desired polarity or molten viscosity profile, these relatively high viscosity polymers may only be employed at minimal concentrations, typically in an amount less than about 40 wt-%, preferably from about 5 wt-% to about 30 wt-%, and more preferably from about 10 wt-% to about 25 wt-%, with respect to the total weight of the composition.

Accordingly, a wide variety of thermoplastic polymers are suitable for use in the present invention. Such thermoplastic polymers are preferably water insensitive. Exemplary polymers for use in the invention include styrenic block copolymers, amorphous and crystalline polyolefins including homogeneous and substantially linear ethylene/alpha-olefin interpolymers; interpolymers of ethylene such as ethylene-vinyl-acetate (EVA), ethylene-methyl acrylate (EMA) and ethylene n-butyl acrylate (EnBa); and mixtures thereof.

A wide variety of block copolymers are useful in the present invention including A-B-A triblock structures, A-B diblock structures, $(A-B)_n$ radial block copolymer structures, as well as branched and grafted versions of such, wherein the A blocks are non elastomeric polymer blocks, typically comprising polystyrene, and the B blocks are unsaturated conjugated diene or hydrogenated version thereof. In general, the B block is typically isoprene, butadiene, ethylene/butylene (hydrogenated butadiene), ethylene/propylene (hydrogenated isoprene), and mixtures thereof. Commercial embodiments include the Kraton® D and G series block copolymers, available from Shell Chemical Company (Houston, TX), Europrene® Sol T block copolymers available from EniChem (Houston, TX), Vector® block copolymers available from Exxon (Dexco) (Houston, TX), as well as others. Block copolymer based compositions are particularly useful for pressure sensitive adhesive applications which generally employ a relatively low melt index block copolymer (less than 50 g/10min) in combination with at least one tackifying resin and plasticizing oil.

Amorphous polyolefins or amorphous polyalphaolefins (APAO) are homopolymers, copolymers, and terpolymers of C_2 - C_8 alphaolefins. These materials are typically polymerized by means of processes which employ Ziegler-Natta catalysts resulting in a relatively broad molecular weight distribution. Commercially available
5 amorphous polyalphaolefins include Rextac® and REXFlex® propylene based homopolymers, ethylene-propylene copolymers and butene-propylene copolymers available from Rexene (Dallas, TX) as well as Vestoplast® alpha-olefin copolymers available from Hüls (Piscataway, NJ).

Metallocene polyolefins are homogeneous linear and substantially linear ethylene
10 polymers prepared using single-site or metallocene catalysts. Homogeneous ethylene polymers are characterized as having a narrow molecular weight distribution and a uniform short-chain branching distribution. In the case of substantially linear ethylene polymers, such homogeneous ethylene polymers are further characterized as having long chain branching. Substantially linear ethylene polymers are commercially available from
15 The Dow Chemical Company as Affinity™ polyolefin plastomers, which are produced using Dow's Insite™ technology, whereas homogeneous linear ethylene polymers are available from Exxon Chemical Company under the tradename Exact®. Homogeneous linear and substantially linear ethylene polymers having a relatively low density, ranging from about .855 to about .885, and a relatively low melt index, for example less than
20 about 50 g/10min. are most preferred, particularly for creating elastomeric fibers, films and adhesive compositions that swell upon exposure to water.

The term "interpolymer" is used herein to indicate a copolymer, terpolymer, or higher order polymer. That is, at least one other comonomer is polymerized with ethylene to make the interpolymer. Interpolymers of ethylene are those polymers having
25 at least one comonomer selected from the group consisting of vinyl esters of a saturated carboxylic acid wherein the acid moiety has up to 4 carbon atoms, unsaturated mono- or dicarboxylic acids of 3 to 5 carbon atoms, a salt of the unsaturated acid, esters of the unsaturated acid derived from an alcohol having 1 to 8 carbon atoms, and mixtures thereof.

30 If employed uncompounded, the ethylene to unsaturated carboxylic comonomer weight ratio is preferably greater than about 3:1, more preferably about 2:1. Hence the

comonomer concentration is preferably greater than 30 wt-%, more preferably greater than 33 wt-% and most preferably greater than 35 wt-%, with respect to the total weight of the ethylene interpolymers. The melt index of the interpolymers of ethylene may range from about 50 to about 2000, preferably from about 100 to 1500, more preferably from about 200 to 1200, and most preferably from about 400 to 1200 g/10 min. When
5 employing a polymer having too low of a melt index uncompounded, the strength of the polymer tends to constrain the swelling of the SAP particles. However, as previously discussed, the disadvantages of the lower melt index interpolymers of ethylene can be overcome by formulating the polymer with diluents.

10 Suitable ethylene/unsaturated carboxylic acid, salt and ester interpolymers include ethylene/vinyl acetate (EVA) ethylene/acrylic acid (EEA) and its ionomers; ethylene/methacrylic acid and its ionomers; ethylene/methyl acrylate (EMA); ethylene/ethyl acrylate; ethylene/n-butyl acrylate (EnBA); as well as various derivatives thereof that incorporate two or more comonomers.

15 Other suitable thermoplastic polymers that may be employed include polylactide, caprolactone polymers, and poly (hydroxy-butyrate/hydroxyvalerate), certain polyvinyl alcohols, biodegradable copolyesters such as Eastman Copolyester 14766 (Eastman), linear saturated polyesters such as Dynapol or Dynacoll polymers from Hüls, poly (ethylene oxide)polyether amide and polyester ether block copolymers available from
20 Atochem (PeBax) or Hoechst Celanese (Rite-flex) respectively, and polyamide polymers such as those available from Union Carbide (Unirez) or Hüls (Vestamelt) or EMS-Chemie (Griltex).

Alternatively, water soluble or water dispersible polymer may be employed provided such polymers are combined with polar diluents such as tackifiers, plasticizers
25 and waxes such that the water sensitivity of the thermoplastic component as a whole is sufficiently reduced such that the composition exhibits the desired rate of acquisition. Representative water soluble or water dispersible polymers that may be employed in this manner include water soluble polyamides such as those described in U.S. Patent No. 3,882,090, incorporated herein by reference, amorphous water sensitive thermoplastic
30 polymers such as polyvinyl alcohol (PVOH) available from Nippon Gohsei as Grohseran L-301 and Grohseran L-302, polyvinyl pyrrolidone (PVP) available from

BASF and ISP, polyvinyl pyrrolidone/vinyl acetate copolymer (PVP/VA) available from ISP, polyetheroxazoline available from Dow under the tradename PEOX and from PCI Incorporated under the tradename Aquazol, polyvinyl methyl ether available from Amoco Chemical Co. under the tradename Amobond, linear polyesters, polyvinyl alcohol, polyacrylamide, water dispersible copolyesters (Eastman AQ).

The thermoplastic component of the composition of the present invention preferably further comprises at least one additional ingredient in an amount up to about 50 wt-%, and preferably from about 10 wt-% to about 40 wt-% with respect to the total composition. Preferred thermoplastic ingredients include those that are commonly employed in hot melt adhesive compositions including plasticizers, tackifiers, waxes, and additives such as antioxidants and pigments. In general, the type of thermoplastic ingredient(s) will be selected to insure sufficient compatibility of the thermoplastic component as a whole. In a preferred embodiment, the thermoplastic component of the composition of the present invention comprises at least one diluent having polar functionality. The diluent is preferably a plasticizer or wax having a molecular weight (Mw) of less than 3000 and preferably less than 2000. The diluent is preferably water insensitive, yet sufficiently polar to reduce the surface tension and/or contact angle of the thermoplastic composition relative to a comparative composition comprising the same ingredients in the absence of such ingredient. Polar diluents include plasticizers and waxes having at least one polar group per molecule. The polar group may be a terminal group or bonded to one or more units in the middle of the molecule. Polar groups include alcohol, ether, ester, epoxy, carboxylic acid, amine, amide, aldehyde, ketone, imide oxime, sulphonic acid, and sulfonamide groups. It is surmised that the polar diluent may plasticize the SAP. Hence, combining the SAP with polar plasticizer alone may increase the absorption rate. However, in the absence of a polymer, plasticizer alone typically could not contribute a continuous thermoplastic phase to disperse the SAP within. Preferably, the diluent is employed at an amount ranging from about 5 wt-% to about 30-wt-% and more preferably in an amount ranging from about 10 wt-% to about 20 wt-% of the total composition.

Exemplary polar plasticizers include phthalate plasticizers such as dioctyl phthalate and butyl benzyl phthalate (e.g., Santicizer 160 from Monsanto); liquid

polyesters such as Dynacol 720 from Huls and liquid polymeric plasticizer available from CP. Hall; benzoate plasticizers such as 1,4- cyclohexane dimethanol dibenzoate (e.g., Benzoflez 352 from Velsicol), diethylene glycol/dipropylene glycol dibenzoate (e.g., Benzoflez 50 from Velsicol), and diethylene glycol dibenzoate where the mole
5 fraction of hydroxyl groups which have been esterified ranges from 0.5 to 0.95 (e.g., Benzoflex 2-45 High Hydroxyl also from Velsicol); phosphite plasticizers such as t-butyl diphenyl phosphate (e.g., Santicizer 154 from Monsanto); liquid rosin derivatives having Ring and Ball softening points below about 60°C such as methyl ester of hydrogenated rosin (e.g., Hercocyn D from Hercules); as well as vegetable and animal oils such as
10 glycerol esters of fatty acids and polymerizable products thereof. Preferred plasticizers include esters of citric acid such as Citroflex® 2, A-2, 4, A-4, A-6, and B-6; butyl benzyl phthalate, toluene sulfonamide, benzoate plasticizers such as 1,4- cyclohexane dimethanol dibenzoate, diethylene glycol/dipropylene glycol dibenzoate, and diethylene glycol dibenzoate where the mole fraction of hydroxyl groups which have been esterified
15 ranges from 0.5 to 0.95.

Further, water soluble or water dispersible plasticizers may also be employed, provided the presence thereof does not impede the rate of absorption of the SAP. Suitable examples include polyethylene glycol with molecular weight below about 2000 and preferably less than 1000, derivatives of polyethylene glycol including Pycal 94, the
20 phenyl ether of PEG available from ICI; ethoxylated bis phenol A (e.g., Macol 206 EM from PPG Industries) and dionyl phenol ethyloxylates (e.g., Surfonic DNP from Huntsman Chemical Corp.).

Exemplary polar waxes include 12-hydroxystearamide, N-(2-hydroxy ethyl 12-hydroxy stearamide (Paricin 220 and 285 from CasChem), stearamide (Kemamide S
25 from Witco), glycerin monostearate, sorbitan monostearate, and 12-hydroxy stearic acid. Also useful in combination with the above are less polar waxes such as N,N'-ethylene-bis stearamide (Kemamide W-40 from Witco), hydrogenated castor oil (castor wax), oxidized synthetic waxes, and functionalized waxes such as oxidized polyethylene waxes (Petrolite E-1040).

30 Other useful plasticizers that may be employed include hydrocarbon oils, polybutene, liquid tackifying resins and liquid elastomers. Plasticizer oils are primarily

hydrocarbon oils which are low in aromatic content and which are paraffinic or naphthenic in character. Plasticizer oils are preferably low in volatility, transparent and have as little color and odor as possible. The use of plasticizers in this invention also contemplates the use of olefin oligomers, low molecular weight polymers, vegetable oils and their derivatives and similar plasticizing liquids.

Waxes are usefully employed to reduce viscosity as well as increase the blocking resistance at concentrations ranging from about 2 wt-% to about 25 wt-% and preferably from about 10 wt-% to about 20 wt-%. Larger concentrations of waxes are typically avoided since waxes tend to bloom to the surface during cooling of the thermoplastic component creating a fluid impermeable barrier layer at the surface of the composition or encapsulating the SAP, thus hindering the ability of the SAP to absorb fluid. In addition to the preferred polar waxes, other useful waxes include paraffin waxes, microcrystalline waxes, Fischer-Tropsch, polyethylene and by-products of polyethylene.

As used herein, the term "tackifier" means any of the compositions described below that are useful to impart tack to the hot melt adhesive composition. ASTM D-1878-61T defines tack as "the property of a material which enables it to form a bond of measurable strength immediately on contact with another surface".

The composition of the invention may comprise up to about 50 wt-% of a tackifying resin. Tackifying resins are preferably employed at a concentration ranging from about 5 wt-% to about 40 wt-% and more preferably from about 10 wt-% to about 20 wt-% with respect to the total weight of the composition.

Tackifying resins comprise resins derived from renewable resources such as rosin derivatives including wood rosin, tall oil and gum rosin as well as rosin esters, natural and synthetic terpenes and derivatives of such. Aliphatic, aromatic or mixed aliphatic-aromatic petroleum based tackifiers are also useful in the invention. Representative examples of useful hydrocarbon resins include alpha-methyl styrene resins, branched and unbranched C₅ resins, C₉ resins and C₁₀ resins, as well as styrenic and hydrogenated modifications of such. Tackifying resins range from being a liquid at 37°C to having a ring and ball softening point of about 135°C.

As is known in the art, various other components can be added to modify the tack, color, odor, etc., of the thermoplastic composition. Additives including

antioxidants such as hindered phenolics (e.g., Irganox™ 1010, Irganox™ 1076), phosphites (e.g., Irgafos™ 168), antiblock additives, pigments and fillers, can also be included in the formulations.

Hydrophilic fillers are a preferred class of additives, which are useful to alter the surface properties and/or increase the rate of absorption. Hydrophilic fillers include calcium carbonate, hydroxyethyl cellulose, hydroxypropyl cellulose and starch or cellulose esters, particularly the acetates.

Surfactants are another preferred additive that reduces the surface tension and/or contact angle of the thermoplastic component. Surfactants are useful in amounts ranging from about 0.5 wt-% to about 25 wt-% and preferably from about 5 wt-% to about 15 wt-%, with respect to the total weight of the thermoplastic component. Suitable surfactants include nonionic, anionic, and silicone surfactants. Exemplary nonionic surfactants are: Ethoxylates of (i) C₁-C₁₈, preferred C₈-C₉ alkyl or dialkyl phenols, such as those sold under the tradenames Macol DNP-10, available from PPG Industries, Gurnee, Ill., a 10 mole ethoxylate of dinonyl phenol, and Triton X-100, available from Union Carbide, a 10 mole ethoxylate of octyl phenol; (ii) alkyl C₈-C₆₀ mono-alcohols, such as those sold under the tradenames Surfonic L-12-8, an 8 mole ethoxylate of dodecanol, available from Huntsman Chemical Co., and Unithox 480, a 38 mole ethoxylate crystalline surfactant available from Petrolite Specialty Polymers Group, Tulsa, Okla.; and (iii) propylene oxide polymers, such as those sold under the tradename Pluronic, which are ethylene oxide/propylene oxide block copolymers having a Mn of 200 to 3000, available from BASF; and benzoates formed by partial condensation of benzoic acid with hydrophilic di or mono-ols having less than 1000 Mn, such as the product of condensing about three equivalents of benzoic acid with four equivalent of diethylene glycol, commercially available as XP 1010 from Velsicol Chemical. A preferred nonionic surfactant blend is Atmer 685, available from ICI Surfactants (Wilmington, DE).

Suitable anionic surfactants are: C₈-C₆₀ alkyl ethoxylate sulfonates, (CH₃-(CH₂)₁₁₋₁₄-(O-CH₂CH₂)₃-SO₃-Na)⁺, such as, Avenel S30, available from PPG Industries; alkyl C₈-C₆₀ sulfonates, such as, Rhodapon UB (C₁₂-SO₃-Na)⁺ available from

Rhone Poulenc; and alkyl/aromatic sulfonates, such as those sold under the tradename Calsoft.

Suitable silicone surfactants are ethoxylates or propoxylates of polydimethyl siloxane, having a number average molecular weight of 500 to 10,000, preferably 600 to 6000, such as are sold under the tradenames Silwet L-77, L-7605, and L-7500 available from OSi Specialties, Danbury, Conn.; and product 193 from Dow Corning.

The preferred surfactants are those with lower molecular weights because these have increased compatibility in the adhesive formulations. The maximum acceptable molecular weight depends on the type of surfactant and the other ingredients in the adhesive formulation.

The thermoplastic component is characterized as having a Brookfield molten viscosity of less than about 30,000 cPs at 177°C, preferably less than about 20,000 cPs, more preferably less than about 10,000 cPs, and even more preferably less than about 5,000 cPs. This viscosity constraint does not necessarily have a lower limit. However, the majority of thermoplastic compositions contemplated for use generally have a viscosity of at least about 50 cPs at 100°C, typically at least about 100 cPs, and preferably at least about 200 cPs. The viscosity is not only critical to the application techniques intended for such composition, but is also critical to the efficiency of the SAP. The present applicants have surprisingly discovered that relatively low viscosity thermoplastic compositions constrain the swelling of the SAP particle to a much lesser extent and hence improve the absorption of the SAP with respect to a high viscosity based composition.

The thermoplastic composition of the present invention comprises at least one SAP polymer admixed with the thermoplastic component. The superabsorbent polymer is present in the composition in an amount ranging from about 5 wt-% to about 70 wt-%, preferably from about 35 wt-% to about 60 wt-%, more preferably ranging from about 40 wt-% to about 60 wt-%, and most preferably from about 50 wt-% to about 60 wt-%. The Applicants have found that by employing SAP at concentration greater than 30 wt-%, particularly in combination with small particle size SAP, a matrix of SAP is formed with the thermoplastic component. The applicants surmise that the formation of this SAP

matrix is essential to the absorption rate of the SAP being unaffected by the presence of the thermoplastic component.

Superabsorbent polymers (SAP), also referred to as water-insoluble absorbent hydrogel-forming polymers (WAHPs) "hydrogel-forming" polymers, and
5 "hydrocolloids" are known and include polysaccharides such as carboxymethyl starch, carboxymethyl cellulose, and hydroxypropyl cellulose; nonionic types such as polyvinyl alcohol, and polyvinyl ethers; cationic types such as polyvinyl pyridine, polyvinyl morpholinone, and N, N-dimethylaminoethyl or N,N-diethylaminopropyl acrylates and methacrylates, and the respective quaternary salts thereof. Typically, WAHPs useful in
10 the present invention have a plurality of anionic, functional groups, such as sulfonic acid, and more typically carboxy, groups. Examples of polymers suitable for use herein include those which are prepared from polymerizable, unsaturated, acid-containing monomers. Such monomers include the olefinically unsaturated acids and anhydrides that contain at least one carbon to carbon olefinic double bond. More specifically, these
15 monomers can be selected from olefinically unsaturated carboxylic acids and acid anhydrides, olefinically unsaturated sulfonic acids and mixtures thereof.

Some non-acid monomers can also be included, preferably in minor amounts, in preparing the SAPs herein. Such non-acid monomers can include, for example, the water-soluble or water-dispersible esters of the acid-containing monomers, as well as
20 monomers that contain no carboxylic or sulfonic acid groups at all. Optional non-acid monomers can thus include monomers containing the following types of functional groups: carboxylic acid or sulfonic acid esters, hydroxyl groups, amide-groups, amino groups, nitrile groups, quaternary ammonium salt groups and aryl groups (e.g., phenyl groups, such as those derived from styrene monomer). These non-acid monomers are
25 well-known materials and are described in greater detail, for example, in U.S. Patent 4,076,663 (Masuda et. al), issued February 28, 1978, and in U.S. Patent 4,062,817 (Westerman), issued December 13, 1977.

Olefinically unsaturated carboxylic acid and carboxylic acid anhydride monomers include the acrylic acids typified by acrylic acid itself, methacrylic acid, ethacrylic acid, -
30 chloroacrylic acid, -cyanoacrylic acid, -methacrylic acid (crotonic acid), -phenylacrylic acid, -acrytoxypionic acid, sorbic acid, -chlorosorbic acid, angelic acid, cinnamic

acid, p-chlorocinnamic acid, -stearylacrylic acid, itaconic acid, citraconic acid, mesaconic acid, glutaconic acid, aconitic acid, maleic acid, fumaric acid, tricarboxyethylene and maleic acid anhydride.

Olefinically unsaturated sulfonic acid monomers include aliphatic or aromatic vinyl sulfonic acids such as vinylsulfonic acid, allyl sulfonic acid, vinyl toluene sulfonic acid and styrene sulfonic acid; acrylic and methacrylic sulfonic acid such as sulfoethyl acrylate, sulfoethyl methacrylate, sulfopropyl acrylate, sulfopropyl methacrylate, 2-hydroxy-3-methacryloxypropyl sulfonic acid and 2-acrylamide-2-methylpropane sulfonic acid.

Other SAPs for use in the present invention contain carboxy groups. These polymers include hydrolyzed starch-acrylonitrile graft copolymers, partially neutralized hydrolyzed starch-acrylonitrile graft copolymers, starch-acrylic acid graft copolymers, partially neutralized starch-acrylic acid graft copolymers, saponified vinyl acetate-acrylic ester copolymers, hydrolyzed acrylonitrile or acrylamide copolymers, slightly network crosslinked polymers of any of the foregoing copolymers, partially neutralized polyacrylic acid, and slightly network crosslinked polymers of partially neutralized polyacrylic acid. These polymers can be used either solely or in the form of a mixture of two or more different polymers. Examples of these polymer materials are disclosed in U.S. Patent 4,076,663 (Masuda et. al), issued February 28, 1978, U.S. Patent 4,093,776 (Aoki et. al), issued June 6, 1978, U.S. Patent 4,666,983 (Tsubakimoto et. al), issued May 19, 1987, and U.S. Patent 4,734,478 (Tsubakimoto et. al), issued March 29, 1988.

Polymer materials used in making the SAPs are slightly network crosslinked polymers of partially neutralized polyacrylic acids and starch derivatives thereof. More preferably still, the SAPs comprise from about 50% to about 95%, more preferably about 75%, neutralized, slightly network crosslinked, polyacrylic acid (i.e., poly (sodium acrylate/acrylic acid)). Network crosslinking renders the polymer substantially water-insoluble and, in part, determines the absorptive capacity and extractable polymer content characteristics of the SAPs. Processes for network crosslinking these polymers and typical network crosslinking agents are described in greater detail in U.S. Patent 4,076,663 (Masuda et. al), issued February 28, 1978.

While the SAP is preferably of one type (i.e., homogeneous), mixtures of polymers can also be used in the present invention. For example, mixtures of starch-acrylic acid graft copolymers and slightly network crosslinked polymers of partially neutralized polyacrylic acid can be used in the present invention.

5 The absorbent gelling particles used in the present invention can have a size, shape and/or morphology varying over a wide range. The absorbent gelling particles may have a large ratio of greatest dimension to smallest dimension (e.g., granules, flakes, pulveruents, interparticle aggregates, interparticle crosslinked aggregates, and the like) and can be in the form of fibers, foams and the like.

10 Particle size is defined as the dimension determined by sieve size analysis. Thus, for example, a particle that is retained on a U.S.A. Standard Testing Sieve with 710 micron openings (e.g., No. 25 U.S. Series Alternate Sieve Designation) is considered to have a size greater than 710 microns; a particle that passes through a sieve with 710 micron openings and is retained on a sieve with 500 micron openings (e.g., No. 35 U.S,
15 Series Aftemate Sieve Designation) is considered to have a particle size between 500 and 710 microns; and a particle that passes through a sieve with 500 micron openings is considered to have a size less than 500.

For particles of SAPs employed in the present invention, the particle size ranges from about 10 to about 1000 microns. Smaller particle sizes, particularly at
20 concentrations greater than 30 wt-%, and preferably about 40 wt-%, have been found to be particularly useful for creating a SAP matrix within the thermoplastic component. Minimal particle size is also advantageous to obtain homogeneous mixing as well as reduce the abrasion and nozzle clogging problems associated with processing relatively large particles using hot melt adhesive application equipment. For these reasons the
25 particle size of the SAP is preferably less than about 500 microns, more preferably less than about 300 microns, and more preferably less than about 200 microns. However, larger particle size may also be employed particularly in applications in which the SAP particle is equal to or larger than the thickness of the thermoplastic component or when an extruder is employed to mix and/or apply the composition.

30 The composition of the present invention is preferably made by first preparing the thermoplastic component by melting and blending all the thermoplastic ingredients and

at 350°F (177°C), and most preferably exhibits the same molten viscosity properties as previously described for the thermoplastic component alone.

The thermoplastic composition of the present invention gels at a rate equal to or faster than that of the superabsorbent polymer alone. Preferably, the composition gels in less than 3 hours, preferably in less than 2 hours, more preferably in less than 1 hour, even more preferably in less than 30 minutes, and most preferably in less than about 15 minutes. Further, for small particle size SAP, the total absorption capacity of the inventive composition is equal to or greater than that of the superabsorbent polymer alone.

The composition of the present invention may be applied by any hot melt application technique such as slot coating, spiral spraying, screen printing, foaming, engraved roller or meltblown adhesive application techniques. When applied in this manner, the inventive thermoplastic composition may be present as a coating, fiber, nonwoven web, or film layer on at least one substrate or as a portion of an article. Preferably, to improve the rate of absorption of the SAP, the application conditions are chosen to maximize the surface areas of the composition. The composition of the present invention may also be formed into a film layer. If a continuous film layer is desired, it is preferred to use a non-contact slot coating application technique described in U.S. Patent No. 5,827,252 issued Oct. 27, 1998, incorporated herein by reference.

The SAP containing thermoplastic composition of the present invention is useful for a variety of end-uses, particularly those which employ superabsorbent polymers such as disposable absorbent articles such as disposable diapers, feminine napkins and medical dressings as well as moisture barriers for optical cable applications and a variety of agricultural applications targeted at increasing humectancy. The inventive composition has further utility in packaging application for food and drugs for absorbing moisture and fluid, for building materials for preventing condensation and waterproofing. Accordingly, the thermoplastic composition may be applied to a variety of substrates using any suitable method, particularly the hot melt adhesive application techniques described above. Substrates in which SAP is commonly applied to includes various film materials such as those employed as diaper backsheets and tape backings, absorbents

then adding the SAP to the molten thermoplastic component. The SAP containing thermoplastic composition may be pelletized, pillowed, or cast into molds or drums, etc., for subsequent remelting and application. Alternatively, all the ingredients may be fed simultaneously at the appropriate rates into an extruder. In the case of extruders, the compounding and application of the SAP containing composition may be achieved in a single process.

In the case of foaming, it is preferred to first coat the SAP particles with a chemical blowing agent. The coated SAP particle is admixed with the thermoplastic component at a temperature below the activation temperature of the chemical blowing agent to prevent premature unintended foaming. The composition is then formed into a film or coated onto a substrate at a temperature or condition sufficient to activate the chemical blowing agent. The chemical blowing agent is surmised to create a gaseous cavity or void with the thermoplastic component with the SAP particle positioned within the void. Preferably, a sufficient amount of coated SAP is employed such that an open cell foam structure is created.

In another embodiment the SAP particles may be coated with the thermoplastic component in particulate form. The coated SAP particle may then be positioned within an article or on a substrate and then heat activated to adhere the SAP particle in place.

The coating of solid particles is done in a number of ways. Both spray drying and prilling are inexpensive methods for coating of particles. In both these processes the particle to be coated is added to the melt or solution and during atomization the coating is done. Another suitable method called Magnetically Assisted Impact Coating (MAIC), available by Aveka Inc. (Woodbury, MN), coats particle onto particle by means of a peening process. By adding a small coating particle onto a large particle core into an assembly of small oscillating magnets, the small particles are readily coated onto the core particles. A preferred method is a two step process of coating the SAP particle with a binder and then contacting the binder coated particle with a powder as described in U.S. Patent No. 5,688,449 issued Nov. 18, 1997, incorporated herein by reference.

The viscosity of the mixture of thermoplastic component and SAP is less than about 200,000 cPs, preferably less than about 100,000, more preferably less than 50,000

such as wood pulp, paper, tissues and towels, nonwovens, and various cable components such as sheathing and jacketing materials.

The invention is further illustrated by the following non-limiting examples. All composition exemplified are expressed in wt-% unless noted otherwise.

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Examples:**Test Methods:**

1. Melt Viscosity is determined in accordance with the following procedure using a Brookfield Laboratories DVII+ Viscometer in disposable aluminum sample chambers. The spindle used is a SC-27 hot-melt spindle, suitable for measuring viscosities in the range of from 10 to 100,000 centipoise. The sample is placed in the chamber, which is in turn inserted into a Brookfield Thermosel and locked into place. The sample chamber has a notch on the bottom that fits the bottom of the Brookfield Thermosel to ensure that the chamber is not allowed to turn when the spindle is inserted and spinning. The sample is heated to the desired temperature, with additional sample being added until the melted sample is about 1 inch (2.5 cm) below the top of the sample chamber. The viscometer apparatus is lowered and the spindle submerged into the sample chamber. Lowering is continued until brackets on the viscometer align on the Thermosel. The viscometer is turned on, and set to a shear rate which leads to a torque reading in the range of 30 to 60 percent. Readings are taken every minute for about 15 minutes, or until the values stabilize, which final reading is recorded.
2. Total Absorption is determined by drawing-down a molten film or heat-pressing the SAP comprising thermoplastic composition into a film. The film is then cut into a 1 inch (2.54 cm) square and weighed. The total amount of SAP in the film is calculated on the basis of the concentration of the SAP within the composition. The sample is then placed in a 25 ml cup and 10 mls of water is poured on top of the film. After 3 hours, the unabsorbed water is filtered off and weighed. The

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total amount of water absorbed is reported as well as the amount of water absorbed per gram of SAP (total absorbed/total SAP).

3. Rate of Absorption is determined by drawing-down a molten film or heat-
 pressing the SAP comprising thermoplastic composition into a film. The film is
 then cut into a 1 inch (2.54 cm) square and weighed. The total amount of SAP in
 the film is calculated on the basis of the concentration of the SAP within the
 composition. The sample is then placed in a 25 ml cup and 10 mls of water is
 poured on top of the film. A stopwatch is started and the time it takes for gelation
 to occur reported.
4. Melt Index is reported in accordance with ASTM D-1238, condition 190°C/2.16
 kg (also known as "Condition E").

15 **Description of Ingredients Employed in the Examples:**

<u>Tradename</u>	<u>Generic Description, Supplier (Location)</u>
<u>Thermoplastic Polymer</u>	
20 Affinity EG 8200	5 MI, .870 g/cm ³ substantially linear ethylene/1-octene interpolymer, Dow (Freeport, TX)
Affinity EG 8100	1 MI, .870 g/cm ³ substantially linear ethylene/1-octene interpolymer, Dow (Freeport, TX)
Eastman Copolyester 14766	biodegradable copolyester, Eastman (Kingsport, TN)
25 Eco PLA	polylactide resin, Cargill (Minneapolis, MN)
EVA 25-400	25% vinyl-acetate, 400 MI EVA
EVA 28-400	28% vinyl-acetate, 400 MI EVA
EVA 28-05	28% vinyl-acetate, 5 MI EVA
EVA 28-800	28% vinyl-acetate, 800 MI EVA
30 EVA 28-2500	28% vinyl-acetate, 2500 MI EVA
EVA 33-44	33% vinyl-acetate, 44 MI EVA

	EVA 33-400	33% vinyl-acetate, 400 MI EVA
	HL-6526	polyamide polymer, (H.B. Fuller Company, St. Paul, MN)
	HL-6582	polyamide polymer, (H.B. Fuller Company, St. Paul, MN)
	HM-1580	polyamide polymer, (H.B. Fuller Company, St. Paul, MN)
5	HL-6594	polyamide polymer, (H.B. Fuller Company, St. Paul, MN)
	HL-6108	polyamide polymer, (H.B. Fuller Company, St. Paul, MN)
	Kraton D-1117	40% diblock, 17% styrene, 106 MI, styrene-isoprene-styrene block copolymer (Shell)
10	Kraton G-1657	35% diblock, , 13% styrene, 370 ¹¹ solution viscosity styrene-ethylene/butylene-styrene block copolymer (Shell)

Plasticizers

	Benzoflex 50	dipropylene-diethylene glycol dibenzoate, Velsicol (Rosemont, IL)
15	Benzoflex 9-88	dipropylene glycol dibenzoate, Velsicol (Rosemont, IL)
	Citroflex A4 special NC)	citrate ester from citric acid, Morflex Inc. (Greensboro, NC)
	Epolene C-3	polyethylene wax, Eastman Chemical Co. (Kingsport, TN)
20	Epolene C-16	polyethylene wax, Eastman Chemical Co. (Kingsport, TN)
	Microwax 180	polyethylene wax
	Paraflint H4	Fischer-Tropsch wax, Moore & Munger (Shelton, CT)
	Paraffin 155	155° melt point paraffin wax
	Penznap N-500	500 viscosity naphthenic oil
25	Santicizer 160	1,2 benzene dicarboxylic acid, butyl phenylmethylester, Solution Inc. (St. Louis, Missouri)
	Uniplex 214	N-butyl benzene sulfonamide, Unitex (Houston, TX)

Tackifiers

30	Escorez 5400	100°C hydrogenated dicyclopentadiene resin (Exxon, Houston, TX)
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Foral AX	hydrogenated rosin, Hercules (Wilmington, DE)
Nirez V-2040	118°C terpene phenolic based resin (Arizona, Panama City, FL)
Zonester 100	94°C rosin based resin (Arizona, Panama City, FL)

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Additives

Surfactant JL80-X	alcohol ethoxylate surfactant, Huntsman (Houston, TX)
Irganox 1010	hindered phenol antioxidant, Ciba Geigy (Hawthorne, NY)
Irganox 1076	hindered phenol antioxidant Ciba Geigy (Hawthorne, NY)

10

Superabsorbent Polymers

Aquaparl 1250H	sodium polyacrylate, Mitsubishi Chemical Co. (Tokyo, JP)
Cabloc AP-80HS	lightly crosslinked sodium polyacrylate, Stockhausen (Greensboro, NC)
15 Favor 800	2-propenoic acid polymers with hydroxyl ethenol homopolymer, Stockhausen (Greensboro, NC)

Due to differences in base chemistry as well as particle size, the gel time of commercially available grades of SAP varies widely. The particle size distribution and gel time of the SAP employed in the examples can be characterized as follows in Table I.

"D(10%), D(50%), and D(90%)" refers to the percentage of particle below X microns.

Hence, in the case of ASAP 2000, 10% of the particles have a particle size of less than 12.5 microns, 50% of the particles have a particle size of less than 62.9 microns, and 90% of the particles have a particle size of less than 561.7 microns.

Table I

	Tradename of SAP			Weight (g)	Gel Time
	D(10%)	D(50%)	D(90%)		
Cabloc 80HS	12.5	62.9	118	0.30	9 minutes
Cabloc 80 HS				0.15	25 minutes
ASAP 2000 (Standard)	208.8	395.5	561.7	0.31	1 Min. 20 Sec
Femdry 31				0.31	20 Sec

Since particle size distribution SAP tends to be very disperse, Favor 800 was
 5 separated into fractions with Vortisiv RBF 8 (MM Industries, Salem, OH) metallic sieve
 and separation equipment. The resulting three portions had particles > 425 microns,
 particles 180- 425 microns, and particles < 180 microns.

Examples 1-5 of Table II employ a thermoplastic adhesive composition
 comprising 80 g of Uniplex 214, 304 g of EVA 28-800, 36 g of Surfactant JL-80X and 4
 10 g of Irganox 1076 in combination with Favor 800 super absorbent polymer at the
 indicated concentration and particle size. By comparing Example 1 to Example 4, the
 effect of increasing SAP concentration as a function of absorption rate is exemplified.
 Example 1 and 4 both contain the same particle size SAP, > 425 microns. However, the
 composition comprising 40 wt-% SAP gelled 2.5 times faster than the composition
 15 employing only 30 wt-%. Examples 2 - 4 employ 40 wt-% of the same SAP differing
 only with respect to the particle size of the SAP. These examples demonstrate that the
 fastest rate of gelation is achieved with the smallest particle size. The sample employing
 <180 micron particle size SAP absorbed the fluid 4 times faster than the sample
 employing the 180-425 micron SAP.

Table II

		Rate of Absorption
Example 1	Adhesive + 30 wt-% > 425 microns	1680 seconds
Example 2	Adhesive + 40 wt-% 180-425 microns	900 seconds
Example 3	Adhesive + 40 wt-% < 180 microns	160 seconds
Example 4	Adhesive + 40 wt-% >425 microns	660 seconds
Example 5	Adhesive + 40 wt-% ASAP 2000	>18000 seconds
Example 6	Adhesive + 40 wt-% AP80HS	840 seconds

μm = micron

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Tables III to XI exemplify compositions of the present invention employing a variety of water insensitive thermoplastic polymers. Example 10 of Table III and Examples 69 and 70 of Table XI employ a substantially linear ethylene interpolymer. Examples 9 and 12-14 of Table III and IV as well as Tables V to IX in their entirety
 10 employ an ethylene interpolymer comprising a polar comonomer. Examples 7, 8 and 11 of Table III and Example 71 of Table XI depict compositions wherein the thermoplastic component is based on a biodegradable thermoplastic material.

Example 7 was found to absorb 10 mls of water in 100 seconds, whereas Example 8 was found to exhibit a slower absorption rate of 190 seconds. Additionally,
 15 Example 8 was noted to be very flexible due to the incorporation of the block copolymer.

Table IV exemplifies the rate of absorption for various inventive compositions of the present invention as well as commercially available comparative examples. According to infrared analysis as well as solvent extraction and centrifugation analytical techniques, Comparative A is believed to comprise about 33.8 wt-% of a 33% vinyl
 20 acetate/44 MI EVA, about 47.3 wt-% high molecular weight (about 10,000) polyethylene glycol, and about 18.9 wt-% sodium salt of polyacrylic acid, whereas Comparative B is believed to comprise about 42.6 wt-% of a 28% vinyl acetate/400 MI EVA, about 26.8 wt-% polyoxyethylene stearate, about 29.5 wt-% polyacrylic acid, and about 1.1 wt-% wax. Comparative C represents a composition made in accordance with Doi et al., U.S.
 25 Patent No. 4,977,211. Specifically, 45 wt-% of LV-780 ethylene-vinyl acetate was combined with 15 wt-% of USR E P02P polyethylene and 40 wt-% of Aquapearl 1250H.

The relatively slow absorption rate is surmised to be attributed to the high cohesive strength of the thermoplastic component.

The EVA content of the inventive EVA based compositions of Tables V and VI varies from 20 wt-% to about 70 wt-%. The rate of gelation increases dramatically at
5 SAP concentrations greater than 30 wt-%. Further, by comparing Examples 15 to 16 of Table V, it is apparent that the addition of plasticizer tends to further increase the rate of acquisition. The Applicants surmise that this effect is attributed to the increased polarity contributed by the plasticizer and/or a further reduction in cohesive strength of the thermoplastic component as a result of the dilution.

10 A 0.15 g sample of the superabsorbent polymer Cabloc 80HS alone, in the absence of being combined with a thermoplastic composition in Table I, took 25 minutes to gel. However, Examples 13-15, all containing about the same amount of SAP gel significantly faster, ranging in rate of gelation from about 3 minutes to about 13 minutes. The Applicants surmise that the thermoplastic component separates the individual SAP
15 particles from each other. Accordingly, each SAP particle can absorb fluid and swell independently, essentially unaffected by the surrounding SAP particles. Hence, gel blocking, the phenomena wherein the gelled SAP particles create a barrier and prevent the migration of fluid to the ungelled SAP particles, is reduced, increasing the rate of gelation.

20 Table VII demonstrates the effect of polar comonomer content and melt index on the rate of gelation. As exemplified by Examples 27, 28 and 30 as the comonomer content increases from 25 wt-% to 28 wt-% to 33 wt-%, the rate of gelation decreases from 24 hours, to about 30 minutes to about 15 minutes. Hence, polar comonomer contents of greater than 25 wt-% are preferred, and greater than 28 wt-% most preferred.
25 Further, as exemplified by Examples 29 and 31-34, as the melt index increases, the gel rate decreases. However, regardless of the melt index of the ethylene interpolymers, all these examples exhibit a substantially faster gel rate than Comparative Examples A to C. The Applicants surmise that the relatively low concentration of SAP causes the reduced gel rate of the comparative examples in view of Examples 21 to 24 of Table IV depicting
30 the gel rate of high SAP containing formulations in comparison to low SAP containing formulations.

Table VIII exemplifies compositions of the present invention comprising wax. In comparison to Example 33, a portion of the plasticizer may be replaced with wax resulting in similar gel rates. The use of Paraffin 155 and Microwax 180 results in faster gel rates. However, as depicted by Example 44, this accelerated rate can only be achieved by employing a combination of polar plasticizer and wax, since wax alone results in a slower gel rate.

Examples 45 to 49 of Table IX exemplify compositions of the present invention comprising tackifier. In comparison to Example 33, by replacing a portion of the plasticizer with tackifier the composition exhibits comparable gel rates. Examples 50 to 56 of Table IX as well as Examples 9 and 12 of Tables III exemplify compositions of the present invention comprising surfactant. Example 37 also contains 4.5 wt-% of Surfactant JL80-X. As in the case of tackifiers, a portion of the polar plasticizer may be replaced with surfactant resulting in comparable and in some instances accelerated gel rates.

Table X exemplifies compositions comprising water insensitive polyamides in combination with polar plasticizer as the thermoplastic component. Employing polyamides is advantageous to increase the heat resistance as well as increase the concentration of SAP. Example 68 employs 70 wt-% SAP, yet remains flexible.

Table XI depicts compositions of the present invention wherein the thermoplastic component contributes elastomeric properties. Examples 69 and 70 employ a substantially linear ethylene/alpha-olefin interpolymer, whereas examples 71 and 72 employ saturated block copolymers. Specifically, Preblend A contains 7 wt-% Kraton G-1651, 17 wt-% Kraton G-1650, 75.25 wt-% Kaydol mineral oil and .25 wt-% Irganox 1010. These compositions can advantageously be sprayed to form a flexible web or slot coated to form a flexible film. The film expands several folds in dimension when exposed to water depending on the concentration of SAP. Surprisingly, the film maintains its integrity after swelling.

Table III

Ingredient Tradename	Ex. 7	Ex. 8	Ex. 9	Ex. 10	Ex. 11	Ex. 12	Ex. 13	Ex. 14
Cabloc AP-80HS	50	50.0	49.8	50	54.90	50.10		40
Aquapearl 1250H								
Irganox 1076			0.4				0.5*	0.5*
Eastman	30	31.5						
Copolyester 14766								
Kraton D-1117		8.0				13.63		
Santicizer 160	20	10.5						
Benzoflex 50			9.10					
Benzoflex 9-88				22.72				
Citroflex A4					10.15		18	18
EVA 28-400			36.2					
EVA 28-800						36.28	41.5	41.5
Eco PLA					34.94			
Surfactant JL80-X			4.50					
Affinity EG 8200				27.27				
Total Parts	100.0	100.0	99.96	99.99	99.99	100	60	100.01

*Irganox 1010

Table IV

Example	Dry Sample Wt.	Time Elapsed until Gelation	Viscosity of Thermoplastic Component @ 350°F (177°C) CPs
Ex. 9	0.3716	2' 38"	
Ex. 11	0.2808	3' 30"	870
Ex. 12	0.2837	4' 0"	3065
Ex. 10	0.4500	15' 0"	
Ex. 13		13"	6740
Ex. 14		12"	11,900
Comparative A	0.3218	> 3 hours	
Comparative B	0.3300	> 3 hours	
Comparative C		300 minutes	

Table V

Example	SAP Cabloc 80 HS (wt-%)	EVA 28-800 (wt-%)	Santicizer 160 (wt-%)	TOTAL (%)	Film Thickness Mils	Film Weight* grams	Gel Time minutes
Ex. 15	50.0	30.0	20.0	100.0	14	0.268	3
Ex. 16	50.0	37.5	12.5	100.0	15	0.263	5
Ex. 17	40.0	45.0	15.0	100.0	14	0.264	13
Ex. 18	30.0	52.5	17.5	100.0	16	0.227	480
Ex. 19	20.0	60.0	20.0	100.0	17	0.242	1440
Ex. 20	10.0	67.5	22.5	100.0	18	0.264	> 4320

*Since the composition comprises 50 wt-% SAP, the weight of SAP = "Film Weight"/2

Table VI

Example	EVA 33-400 (wt-%)	EVA 33-44 (wt-%)	Benzoflex 50 (wt-%)	Santicizer 160 (wt-%)	Cabloc 80-HS (wt-%)	Film Thickness mils	Film Weight grams	Gel Time
Ex. 21	11.16	18.84		20.00	50.00	9.0	0.1762	7 min 25 sec
Ex. 22	17.85	30.15		32.00	20.00	10.6	0.1931	over 6 hours
Ex. 23	11.16	18.84	20.00		50.00	11.5	0.19999	8 min & 40 sec
Ex. 24	17.85	30.15	32.00		20.00	11.8	0.1800	over 6 hours

Table VII

Example #	25	26	27	28	29	30	31	32	33	34
Ingredient										
Cabloc 80 HS	50	50	50	50	50	50	50	50	50	50
EVA 25-400			50							
EVA 28-400						50			35	
EVA 33-400				50						
EVA 28-800	36.3	34.5						30		
EVA 28-2500							50			30
EVA 28-03					18.2					
Citroflex A-4		15			31.8			20	15	20
Uniplex 214	13.2									
Irganox 1010	0.5	0.5								
Tests results and observations										
Film appearance	Flexible bleed	Flexible	Strong flexible	Strong flexible	Flexible	Strong flexible	Strong flexible	Flexible	Good flexible film	Flexible
Avg. Gel time (1"x1" (2.54 cm x 2.54 cm) about 0.3g film in 10ml water)	5 min	6 min	over 24 hours	15 min. 30 sec.	Expands while gelling (13 min)	36 min. 15 sec.	6 min 15 sec	4 min. 33 sec.	6 min 50 sec	1 min 40 sec

Table VIII

Example No.	35	36	37	38	39	40	41	42	43	44
Cabloc 80 HS	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0
EVA 28-800	29.7	29.7	29.7	29.7	29.7	29.7	24.0	24.0	24.0	34.5
AC-400	8.0									
Paraffin 155F		8.0					16.0			
Microwax 180F			8.0					16.0		15.0
Epolene C-3				8.0					16.0	
Epolene C-16					8.0					
Parafint H4						8.0				
Citroflex A-4	12.0	12.0	12.0	12.0	12.0	12.0	9.7	9.7	9.7	
Irganox 1010	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.5
Property Tests results and Gel time (fluid absorption)										
Film appearance	less flexible	less flexible	flexible	flexible	flexible	flexible	slightly flexible	slightly brittle	slightly brittle	slightly brittle
Avg. Gel time (1"x1" (2.54 cm x 2.54 cm) about 0.3g film in 10 ml water)	5 minutes	3 minutes 15 seconds	3 minutes 45 seconds	6 minutes 20 seconds	6 minutes 20 seconds	6 minutes 20 seconds	7 minutes 45 seconds	2 minutes 30 seconds	2 minutes 52 seconds	9 minutes 45 seconds

Table IX

Example No.	45	46	47	48	49	50	51	52	53	54	55	56
Ingredient												
Cabloc 80 HS	50.0	50.0	50.0	50.0	50.0	48.9	47.0	47.0	47.0	40.0	40.0	40.0
EVA 28-800	29.7	29.7	29.7	29.7	24.0	33.6	32.4	32.4	32.4	40.0	40.0	41.5
CAB-O-SIL-m5						2.5						
Emerest 2400							6.0					
Pluronic F-38								6.0		5.0		
Pluronic F0108									6.0		5.0	
Zonester 100				8.0	16.0							
Nirez V-2040	8.0											
Escorez 5400		8.0										
Foral AX			8.0									
Citroflex A-4	12.0	12.0	12.0	12.0	9.7	14.5	14.1	14.1	14.1	15.0	15.0	18.0
Irganox 1010	0.3	0.3	0.3	0.3	0.3	0.5	0.5	0.5	0.5			0.5
Property Tests results and Gel time (fluid absorption)												
Film appearance	flexible	flexible	flexible	sets slow, flexible	sets slow, flexible	flexible	flexible	flexible	flexible	flexible	flexible	flexible, sprayable
Avg. Gel time (1"x1" (2.54 cm x 2.54 cm) about 0.3g film in 10 ml water)	5 minutes 40 seconds	6 minutes 30 seconds	7 minutes 25 seconds	5 minutes 15 seconds	5 minutes 20 seconds	6 minutes 30 seconds	6 minutes 40 seconds	6 minutes 35 seconds	5 minutes 5 seconds	4 minutes 35 seconds	4 minutes 50 seconds	

Table X

Example #	57	58	59	60	61	62	63	64	65	66	67	68
Ingredient												
HL-6526	42.0	32.0						8.0	6.4		40	24.0
HL-6582x			42.0	32.0		20.0		32.0	25.6			
HM 1580					40.0	20.0						
HL 6594							40.0					
HL 6108										40.0		
Benzoflex 9-88							10.0					
Uniplex 214	8.0	8.0	8.0	8.0	10.0	10.0		10.0	8.0	10.0	10	6.0
Cabloc 80	50.0	60.0	50.0	60.0	50.0	50.0	50.0	50.0	60.0	50.0	50	70.0
HS												
Irganox 1010												
Test results and observations												
Film appearance	slow setting flexible	slow setting flexible	brittle	very brittle	flexible	flexible	slow setting tacky	brittle	brittle	flexible slow setting and tacky	flexible	flexible
Avg. Gel time (1"x1" (2.54 cm x 2.54 cm) about 0.3g film in 10ml water)	7 min. 5 sec.	5 min.	4 min. 20 sec.	4 minutes	7 min	4 min. 30 sec.	2 min 40 sec.	2 min. 20 sec.	2 min	2 min. 20 sec.	12 minutes 20 seconds	

Table XI

Example No.	69	70	71	72
Ingredient				
Cabloc 80 HS	50	50	50	50
Irganox 1010				
Benzoflex 8-88	12	15		10
Cirolflex A-4				
Penznap N500 oil	26	20		
Kraton G1657		5		
Preblend A			50	40
Copolyester 14766				
EG 8100	12	10		
Property test results				
Film	very elastic, flexible, expanded in fluid bleed.	didn't bleed, flexible, elastic, expandable	flexible	flexible
Gel time	over 30 min.	7 minutes	Expanded in one hr.	7 minutes

What is claimed is:

1. A thermoplastic composition selected from the group consisting of:
 - a) compositions comprising a thermoplastic component comprising at least one
5 thermoplastic polymer and at least one diluent having polar functionality and at least one superabsorbent polymer;
 - b) compositions comprising a thermoplastic component comprising at least one interpolymer of ethylene having a melt index greater than about 100 g/10 minutes and a comonomer content greater than 30 wt-% and at least one superabsorbent polymer;
 - 10 c) compositions comprising about 25 wt-% to about 50 wt-% of a thermoplastic component having a Brookfield viscosity of less than 30,000 cPs at about 180°C and about 35 wt-% to about 75 wt-% of a superabsorbent polymer; and
 - d) compositions comprising about 25 wt-% to about 75 wt-% of a thermoplastic component and about 25 wt-% to about 75 wt-% of at least one superabsorbent
15 polymer wherein the superabsorbent gels in less than 3 hours.
2. The thermoplastic composition of Claim 1 wherein said thermoplastic component is water insensitive.
- 20 3. The thermoplastic composition of Claims 1-2 wherein said diluent is a plasticizer.
4. The thermoplastic composition of Claim 1-2 wherein said diluent is a wax.
5. The thermoplastic composition of Claim 1-4 wherein the diluent is present in an
25 amount ranging from about 5 wt-% to about 30 wt-%.
6. The thermoplastic composition of Claim 1 wherein the melt index of the interpolymer of ethylene is greater than 400 g/10 min.

7. The thermoplastic composition of Claim 1 wherein the melt index of the interpolymers of ethylene is greater than about 800 g/10 min.
8. The thermoplastic composition of Claim 1 wherein the melt index of the interpolymers of ethylene is greater than about 1000 g/10 min.
9. The thermoplastic composition of Claims 1 and 6-8 wherein the comonomer of the interpolymers of ethylene content is greater than 33 wt-%.
10. The thermoplastic composition of Claims 1 and 6-9 wherein the comonomer is selected from the group consisting of vinyl acetate, n-butyl acrylate, methacrylate and mixtures thereof.
11. The thermoplastic composition of Claim 10 wherein the thermoplastic component consists essentially of:
 - a) about 10 wt-% to about 50 wt-% of at least one water insoluble polymer;
 - b) 0 wt-% to about 30 wt-% of a plasticizer having polar functionality; and
 - c) 0 wt-% to about 40 wt-% of a tackifier, wax, non-polar plasticizer, and mixture thereof.
12. The thermoplastic composition of Claims 1 to 11 wherein the superabsorbent polymer is present in an amount ranging from about 30 wt-% to about 60 wt-%.
13. The thermoplastic composition of Claims 1 to 5 wherein the thermoplastic component comprises at least one polymer selected from the group consisting of block copolymers, amorphous polyolefins, crystalline polyolefins, interpolymers of ethylene and mixtures thereof.
14. The thermoplastic composition of Claims 1 to 13 wherein the composition gels at a rate substantially equal to or faster than that of the superabsorbent polymer alone.

15. The thermoplastic composition of Claims 1 to 13 wherein the composition gels in less than 1 hour.
- 5 16. The thermoplastic composition of Claims 1 to 13 wherein the composition gels in less than 30 minutes.
17. The thermoplastic composition of Claims 1 to 16 wherein the thermoplastic component is a hot melt adhesive.
- 10 18. A film layer disposed on a substrate comprising the thermoplastic composition of Claims 1 to 17.
19. The thermoplastic composition of Claims 1 to 17 wherein superabsorbent polymer has a particle size equal to or greater than the thickness of the film layer.
- 15 20. A method of making a film comprising the steps of providing a molten mixture of a composition of Claims 1 to 17 and forming a substantially continuous film from said molten mixture.
- 20 21. A method of applying superabsorbent polymer to a substrate comprising the steps of:
- a) providing a molten mixture comprising the thermoplastic composition of Claims 1 to 17; and
 - b) applying said mixture to a substrate.
- 25 22. The method of Claim 21 wherein said substrate is selected from the group consisting of film, paper, nonwoven, tissue, and an absorbent core.
23. A nonwoven web comprising the composition of Claims 1 to 17.

24. A disposable article comprising a body fluid pervious topsheet and a body fluid impervious backsheet and the composition of Claims 1 to 17 disposed between said topsheet and backsheet.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/10135

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C08L101/00 A61L15/00 H01B7/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61L H01B C08L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 248 709 A (HELMUT BREHM) 28 September 1993 (1993-09-28) see the whole document ---	1-24
X	US 5 567 744 A (MANABU NAGATA ET AL.) 22 October 1996 (1996-10-22) see the whole document ---	1-24
X	US 5 246 770 A (VINCENT BOTTIGLIONE ET AL.) 21 September 1993 (1993-09-21) column 2, line 34 - column 3, line 9 ---	1-24
Y	WO 91 18042 A (THE DOW CHEMICAL COMPANY) 28 November 1991 (1991-11-28) page 6, line 24 - line 31 --- -/--	1-24



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Patent family members are listed in annex.

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Date of the actual completion of the international search

13 August 1999

Date of mailing of the international search report

26/08/1999

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>PATENT ABSTRACTS OF JAPAN vol. 96, no. 5, 31 May 1996 (1996-05-31) & JP 08 027372 A (NIPPON SHOKUBAI), 30 January 1996 (1996-01-30)- abstract</p> <p>-----</p>	1-24

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/10135

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5248709 A	28-09-1993	DE 3741157 A	15-06-1989
		AT 118798 T	15-03-1995
		CA 1338730 A	26-11-1996
		DE 3853144 D	30-03-1995
		WO 8905326 A	15-06-1989
		EP 0321755 A	28-06-1989
		EP 0389547 A	03-10-1990
		GR 90300017 T	07-06-1991
US 5567744 A	22-10-1996	JP 3501494 T	04-04-1991
		JP 5320523 A	03-12-1993
		CA 2136420 A	09-12-1993
		CN 1082429 A	23-02-1994
		EP 0641835 A	08-03-1995
US 5246770 A	21-09-1993	WO 9324575 A	09-12-1993
		FR 2640547 A	22-06-1990
		AT 103730 T	15-04-1994
		CA 2005764 A	20-06-1990
		DE 68914274 D	05-05-1994
		DE 68914274 T	14-07-1994
		EP 0378940 A	25-07-1990
		ES 2057168 T	16-10-1994
		JP 2667266 B	27-10-1997
WO 9118042 A	28-11-1991	JP 3121834 A	23-05-1991
		AU 7863091 A	10-12-1991
		CA 2082623 A	20-11-1991
JP 08027372 A	30-01-1996	EP 0530231 A	10-03-1993
		NONE	